

ACRYLATE-FUNCTIONAL BLOCKED POLYISOCYANATE
RESIN FOR UV/THERMALLY CURABLE COATINGS

BACKGROUND OF THE INVENTION

The present invention relates to a solvent-based, dual cure coating incorporating thermally curable blocked polyisocyanate resins, which also contain unsaturated linkages.

The hardening of coating compositions containing (meth)acryloyl
5 groups by UV-initiated polymerization which proceeds by a free radical mechanism is a known and industrially established procedure. It is one of the most rapid methods of hardening in coating technology. However, UV-hardening lacquers have certain disadvantages, such as their high volume shrinkage during hardening, which can result in adhesion problems. The
10 high crosslink density which is inherent in the system results in brittleness, lack of flexibility and a lack of impact strength. Also, the hardening of coating compositions containing UV-hardening (meth)acryloyl groups is dependent on the availability of a sufficient dosage of radiation of the appropriate wavelength. Poorly exposed regions suffer from reduced
15 hardening, which results in appreciable losses in surface resistance, and unexposed regions do not harden at all.

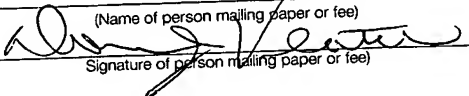
Also well known and widely used in industrial practice is the preparation of coatings from aqueous compositions derived from polyisocyanates and compound with isocyanate-reactive groups. Due to
20 the wide choice of reactants, the technical properties of these coatings, such as adhesion, flexibility, impact strength and resistance to weathering, can be varied over a wide range. However, compared with radiation-hardening systems, the hardening of aqueous two-component polyurethane compositions requires a long time, even at elevated
25 temperatures. Another difference is that UV light is not necessary for curing.

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Combinations of both types of hardening, i.e., so-called dual cure systems hardening by UV radiation and hardening by polyurethane formation, are also known for solvent-based lacquers. For example, U.S. Patent 4,342,793 describes hardenable resin compositions which contain
5 a radiation-hardenable reactive thinner (low molecular weight esters of acrylic acid), a saturated polyol and a polyisocyanate. Hardening is carried out by irradiation in order to polymerize the reactive thinner, which is followed by thermal hardening to form a polyurethane from the polyol and the polyisocyanate. One-component dual cure coatings are also known
10 (see U.S. Patent 5,234,970).

Dual cure systems have varying processing (e.g. processing speed, pot life and cure conditions) and end use demands (e.g., thermal characteristics, weathering and hydrolytic stability). Accordingly the intended application and the environment related to that application have
15 typically dictated the particular formulation of the coating.

U. S. Patent 5,350,823 discloses the preparation of a self-crosslinkable copolymer of a) monomer prepared by first reacting isophorone diisocyanate (IPDI) with 2-hydroxyethyl acrylate and then with butanone oxime, and b) an unsaturated compound. Crosslinking of the
20 copolymer occurs by a single mechanism only.

U.S. Patent 4,485,226 describes a UV-curable composition which is a combination of a) a blocked isocyanate prepared by reacting a portion of the isocyanate groups of an aliphatic isocyanate with butanone oxime and then reacting the rest of the isocyanate groups with hydroxyethylacrylate
25 and b) an isocyanate reactive material (see column 5, lines 19-31). As described therein, after polyurethane formation, the systems are converted to the cross-linked state by high-energy radiation.

U.S. Patent 5,908,907 describes aqueous emulsions from blocked isocyanates wherein a portion of the isocyanate groups of an HDI trimer
30 are blocked with methyl ethyl ketoxime and then the rest of the isocyanate groups are reacted with hydroxyethylacrylate.

U.S. Patent 4,008,247 discloses reacting a portion of the isocyanate groups of an aromatic isocyanate with an hydroxy functional unsaturated monomer and the rest of the isocyanate groups with a blocking agent (including dimethyl pyrazole). Ethylenically unsaturated blocked (cyclo)aliphatic diisocyanates are not described.

There remains a need for dual-cure coatings formulations, producing desirable coatings properties, in which blocked (cyclo)aliphatic polyisocyanates are combined in the same molecule with UV cure capabilities.

DESCRIPTION OF THE INVENTION

As used herein, unless otherwise expressly specified, all of the numerical ranges, amounts, values and percentages such as those for amounts of materials, times and temperatures of reaction, ratios of amounts, values for molecular weight, and others in the following portion of the specification may be read as if prefaced by the word "about" even though the term "about" may not expressly appear with the value, amount or range.

The present invention is directed to a non-aqueous, dual-cure composition, its method of preparation and its method of use. The non-aqueous, dual-cure composition comprises:

- a) from about 5 to about 85% by weight of a blocked (cyclo)aliphatic polyisocyanate prepared by
 - i) reacting a hydroxy-functional (meth)acrylate with an organic (cyclo)aliphatic polyisocyanate at an NCO to OH equivalent ratio of from about 1.5:1 to about 3:1, with the resultant partially blocked isocyanate having an isocyanate group content of from about 5 to about 13% by weight, and

- The blocked (cyclo)aliphatic polyisocyanate (component a)) is prepared by reacting a hydroxy-functional (meth)acrylate with an organic (cyclo)aliphatic polyisocyanate at an NCO to OH equivalent ratio of from about 1.5:1 to about 3:1, preferably from about 1.8:1 to about 2.2:1, with the resultant partially blocked isocyanate having an isocyanate group content of from about 5 to about 13% by weight, preferably from about 8 to about 13% by weight, and reacting the remaining isocyanate groups with a blocking agent. The dual-cure composition contains from about 5% to about 85%, preferably from about 14 to about 75% by weight of the blocked (cyclo)aliphatic polyisocyanate (component a)).

The unsaturated, hydroxy-functional (meth)acrylates useful in preparing the blocked isocyanate include, for example hydroxy-C₁-C₁₀-alkyl (meth)acrylates. Specific examples include, 2-hydroxyethyl acrylate, 2-hydroxypropyl acrylate, 2-hydroxyethyl methacrylate, 3- and 4-hydroxybutyl acrylate, 3- and 4-hydroxybutyl methacrylate, 6-hydroxyethyl acrylate, 6-hydroxyethyl methacrylate, 3-hydroxy-2-ethylhexyl acrylate, 3-hydroxy-2-ethylhexyl methacrylate, and, 2-hydroxypropyl methacrylate. Mixtures can also be used. The use of 2-hydroxyethyl acrylate is preferred.

Substantially any aliphatic and/or cycloaliphatic isocyanate may be used. Examples include organic (cyclo)aliphatic polyisocyanates of the general formula $X(NCO)_2$, where X is an aliphatic hydrocarbon radical of 4 to 12 carbon atoms, or a cycloaliphatic hydrocarbon radical of 6 to 15 carbon atoms, may be used according to the present invention. Specific examples of useful isocyanates include, but are not limited to, butane-1,4-diisocyanate, hexane-1,6-diisocyanate, 2,2,4- and 2,4,4-trimethyl-hexamethylene diisocyanate, cyclohexane diisocyanate, isophorone diisocyanate, 1,4-bis(β -isocyanatoethyl) cyclohexane, isophorone diisocyanate, 4,4'-diisocyanatodicyclohexylmethane and hexamethylene diisocyanate. Also useful are isocyanurate-group containing polyisocyanates based on aliphatic and/or cycloaliphatic isocyanates. The production of isocyanurate-group containing polyisocyanates is known in the art and is described, for example, in U.S. Patents 4,288,586 and 4,324,879. Also useful are uretdione-containing, biuret-containing and urethane-group containing isocyanates based on aliphatic and/or cycloaliphatic isocyanates. Such isocyanates are known in the coating art. The use of a low viscosity solvent-free polyfunctional aliphatic polyisocyanates based on hexamethylene diisocyanate (such as the resin commercially available from Bayer Polymers LLC as Desmodur N 3600) is preferred.

Blocking agents are known in the coatings art. Examples of useful blocking agents include, but are not limited to, acetone oxime, diethyl ketoxime, methyl isopropyl ketoxime, methyl isobutyl ketoxime, diisopropyl ketoxime, methyl ethyl ketoxime, dimethylpyrazole, malonic acid dimethyl ester, acetoacetic acid ethyl ester, ϵ -caprolactam, pyrazole and C₁ to C₄ alkyl pyrazoles. Dimethylpyrazole (DMP) is presently preferred.

In preparing the blocked polyisocyanate (component a)), typically, the (cyclo)aliphatic polyisocyanates are combined with solvent and catalyst, with stirring and heating. The hydroxy-functional (meth)acrylate is added in dropwise fashion to the (cyclo)aliphatic polyisocyanates, with stirring and heating until the desired NCO content is obtained. The temperature is then increased and the blocking agent is added until the NCO peak (as observed through IR spectrophotometry) disappears. The resultant blocked isocyanate contains both blocked isocyanate functionality and ethylenically unsaturated functionality.

The hydroxy-functional polymer (ingredient b)) has an OH number from about 10 to 250. The hydroxy-functional polymer also has an acid number of from about 0.1 to about 50, preferably from about 1 to about 5. The equivalent ratio of blocked isocyanate groups to hydroxy groups in the hydroxy-functional polymer ranges from about 0.8:1 to about 1.2:1, preferably from about 0.9:1 to about 1.1:1. The hydroxy-functional polymer may be a saturated polyester or an unsaturated polyester, or a mixture thereof. Suitable polyesters are known in the polyurethane art. Saturated polyesters are preferred, with linear saturated polyesters being most preferred. The composition contains from about 5% to about 85%, and preferably from about 25% to 86% by weight, of the hydroxy-functional polymer.

The ethylenically unsaturated ingredient (component c)) contains a compound selected from the group consisting of ethylenically unsaturated monomers, polymers containing ethylenic unsaturation (other than unsaturated polyesters) and mixtures thereof. It is preferred that the

ethylenically unsaturated ingredient contains two or more ethylenic double bonds. Preferred as the ethylenically unsaturated component are di-(meth)acrylates of glycols containing from 2 to 6 carbon atoms and poly(meth)acrylates of polyols containing from 3 to 4 hydroxyl groups and 3 to 6 carbon atoms. Specific useful di-(meth)acrylates and poly(meth)acrylates include ethyleneglycol diacrylate, 1,3-propanediol diacrylate, 1,4-butanediol diacrylate, 1,6-hexanediol diacrylate, trimethylolpropane triacrylate, pentaerythritol tetraacrylate, and the corresponding di- and polymethacrylates. Also useful are the di-(meth)acrylates of ether-group containing diols and the poly(meth)acrylates of ether-group containing polyols. The presently preferred unsaturated compound is 1,6-hexanediol diacrylate. The composition contains from 0 to about 65% by weight, and preferably from about 10 to about 60% by weight, of the ethylenically unsaturated ingredient. As noted above, if component b) is a saturated polyester, component c) must be present in an amount of from about 20 to about 65% by weight.

As the UV initiator, any of the compounds known in the art for promoting free radical polymerization using conventional ultraviolet light sources may be used in the coating. UV initiators for free-radical polymerization (component d)) include aromatic ketones such as benzophenone and benzoin ethers such as benzoin isobutyl ether. Other useful photoinitiators are described, e.g., in U.S. Patents 4,485,226 and 5,234,970. The presently preferred UV initiator for free-radical polymerization is 1-hydroxy-cyclohexyl phenyl ketone, commercially available from Ciba Specialty Chemicals as Irgacure® 184. The composition contains from about 0.1% to about 7%, preferably from about 0.5% to about 6% of the UV initiator for free-radical polymerization.

Catalysts for promoting the cure of the reaction between the isocyanate groups (of component a)) and the hydroxy groups (of

component b)) can also be added. Examples include, for example, the dialkyl tin salts of a long chain fatty acid such as dibutyltin dilaurate.

Other additives that can be used include flow and leveling aids, mar and slip agents, stabilizers, UV absorbers, antioxidants, inorganic and organic pigments, fillers (such as quartz powder, finely divided silica, barite, calcium carbonate, chalk, dolomite and talc), pigment dispersants and the like.

Coatings may be prepared by combining ingredients a), b), c), and d), to produce a formulation, applying the formulation to a surface to be coated, and curing the resulting formulation. Two curing processes are used: a baking or thermal cure and a UV cure. The curing processes may be performed in either order. For the thermal cure, the coating is typically heated to a temperature of from about 200 to about 750°F for a period of from about 10 seconds to about 30 minutes. Suitable UV radiation sources are known in the art and include artificial radiation sources emitting in the range of from about 250 to about 500 nm and preferably in the range of from about 300 to about 400nm. Mercury vapor, xenon and tungsten lamps may be used. While the total dose of radiation depends upon the thickness of the film applied, generally it amounts to between from about 2000 to about 3000 mJ/cm².

In the examples, all figures cited in percent or in parts represent percentages by weight and parts by weight, respectively, unless otherwise indicated.

EXAMPLES

25 Example 1: Preparation of Blocked Polyisocyanate Acrylate 1

Into a two liter flask fitted with stirrer, heater, air inlet, distillation column and addition funnel was combined 618.35 g (3.39 eq) of Desmodur N-3600 (a polyfunctional aliphatic polyisocyanate resin based on hexamethylene diisocyanate (HDI), available from Bayer Polymers LLC, having an NCO content of 23.0% by weight) and 240.16 g of N-methyl

pyrrolidone. Air was bubbled below the surface of the liquid, and nitrogen gas purged above the liquid surface. To this, 0.97g of 2,6-di-tert-butyl-4-methylphenol (BHT) and 0.003 g of dibutyltindilaurate was added with stirring and the mixture heated to 60°C. Dropwise addition of 129.52 g (1.11 eq) of 2-hydroxyethylacrylate was made while the temperature was maintained between 60 to 70°C. A reaction temperature of 60°C was maintained until the mixture reached the theoretical NCO content of 9.34%. The temperature was then increased to 100°C, and slow addition of 215.76 g (2.22 eq) of 3,5-dimethylpyrazole was started while the temperature was maintained at 100°C. The reaction was checked for completion by monitoring the disappearance of the NCO peak (2263-2275 cm^{-1}) by IR spectrophotometry. The product was cooled to room temperature and poured into a 1 liter aluminum bullet container sealed under a head of dry air. The resin properties are summarized in Table 1 below:

Table 1

Solids Content	80% by weight
Viscosity, 25°C	12,700 cps
Density	9.4 lbs/gal
Blocked Isocyanate Content	7.74% by weight
Blocked Isocyanate Equivalent Weight	543

20 Example 2: Preparation of Blocked Polyisocyanate Acrylate 2

Into a two liter flask fitted with stirrer, heater, air inlet and distillation column was combined 581.4 g (1.72 eq) of the reaction product of Desmodur N-3600 and hydroxyethylacrylate having an NCO content of 12.8% and a viscosity at 23°C of 12,500 cps and 250.0 g of n-butyl acetate. A nitrogen pad was maintained on top of the condenser while stepwise addition of 168.6 g (1.76 eq) of 3,5-dimethylpyrazole was begun, and while a temperature of less than 50°C was maintained. After addition

of all of the 3,5-dimethylpyrazole, the reaction mixture was maintained at a temperature of 50°C, until the NCO content was less than 0.05% by titration. The product was cooled to room temperature and poured into a 1 liter aluminum bullet container sealed under a head of dry air. The resin properties were as summarized in Table 2 below:

Table 2

Solids Content	75% by weight
Viscosity, 25°C	820 cps
Density	9.0 lbs/gal
Blocked Isocyanate Content	7.23% by weight
Blocked Isocyanate Equivalent Weight	581

10

Example 3 through 10: Formulation and Testing

Formulations prepared according to the present invention were cured and tested. The formulations were prepared to produce a final NCO/OH ratio of 1.0:1.0. A Byrd applicator draw down bar with a 4 mil gap was used to apply the coating to glass. The dry film thickness was from 1.5 to 2 mils. The cure was achieved by two different procedures. In one case (A), the UV cure was done first followed by oven bake; in the other case (B), the oven bake was done first, followed by UV cure. Cure schedule (A) called for UV curing (10 fpm\100% power using Fusion UV Systems conveyor unit and "H"-bulbs) followed by baking at 260°F for 30 minutes. Cure schedule (B) called for baking at 260°F for 30 minutes followed by UV curing (10 fpm\100% power using Fusion UV Systems conveyor unit and "H"-bulbs). The UV curing accounted for approximately 2000 – 3000 mJ/cm².

25

Examples 3, 4, 7 and 8 are comparative examples (since the hydroxy-functional polymer was a saturated polyester and no ethylenically unsaturated compound was added). The formulations used were as set forth in Tables 3 and 4, and the materials were as follows:

5

Polyester A - a saturated polyester prepared from phthalic acid, adipic acid and trimethylolpropane. Polyester A was used as 75% by weight solids in propylene glycol monomethyl ether acetate. The polyester has an OH number of 210 and an acid number of 3.

10

Polyester B - a linear, saturated polyester prepared from adipic acid, ethylene glycol and diethylene glycol. The polyester has an OH number of 56 and an acid number of 2.

15

1,6-HDDA - 1,6-hexanediol diacrylate

IR-184 - Irgacure 184 - 1-hydroxycyclohexylphenyl ketone, available from Ciba Specialty Chemicals

20

BLA1 - the blocked polyisocyanate acrylate of Example 1

Table 3

		3	4	5	6
Formulations	Polyester A, pbw	50		50	
	Polyester B, pbw		50		50
	1,6-HDDA, pbw			54	54
	IR-184, pbw	2.67	2.05	4.65	3.66
	BLA1, pbw	68.59	18.3	68.59	18.3
	BA, pbw	20	10		

Table 4

		7	8	9	10
Formulations	Polyester A, pbw	50		50	
	Polyester B, pbw		50		50
	1,6-HDDA, pbw			54	54
	IR-184,pbw	5.34	4.1	9.3	7.32
	BLA1, pbw	68.59	18.3	68.59	18.3
	BA, pbw	20	10		

The coatings were tested for chemical resistance and hardness. The chemical resistance was a methylethyl ketone resistance test. This test was performed using an 8"x8" cheesecloth and a 2-pound ball peen hammer. The dry film thickness of the coating was measured and the coated panel was wiped off with a cloth to remove any loose materials. The coated panel was then placed on a flat surface with the painted side up. The ball peen was securely wrapped with several layers of cloth (8"x8" cloth folded twice). The cloth was secured to the hammer by using a rubber band to insure that the cloth did not come off during test. The cheesecloth was saturated with methylethyl ketone. The wet ball peen was placed on the coated surface, with the hammer being held so that the ball peen was at a 90-degree angle to the panel. Without applying downward pressure, the hammer was moved back and forth over an approximately 4" long area of the coating. One forward and back motion was counted as 1 double rub. The cloth was saturated with MEK after every 25 double rubs. The test was stopped if breakthrough of the coating occurred, with the number of double rubs being recorded. Hardness was tested according to ASTM-D4366-95. The results of both tests were as set forth in Table 5. In the Table, DR represents the number of double rubs recorded and PI representing the initial pendulum hardness in seconds.

Table 5

Example #		3	4	5	6	7	8	9	10
Cure schedule									
A	DR	21	2	>100	41	40	6	>100	30
B	DR	30	30	80	45	46	10	62	13
A	PI	101	19 tacky	122	22 tacky	106	27 tacky	75	21 tacky
B	PI	40	18 tacky	50	37	62	15 tacky	67	34

Although the invention has been described in detail in the foregoing for the purpose of illustration, it is to be understood that such detail is solely for that purpose and that variations can be made therein by those skilled in the art without departing from the spirit and scope of the invention except as it may be limited by the claims.